



wherein o is 0-6 and wherein the phenyl radical is optionally substituted up to twice, each substituent chosen independently from F, Cl, Br, OH, CF<sub>3</sub>, NO<sub>2</sub>, CN, OCF<sub>3</sub>, (C<sub>1</sub>-C<sub>6</sub>)-alkoxy, (C<sub>1</sub>-C<sub>6</sub>)-alkyl, and NH<sub>2</sub>; NH<sub>2</sub>, NH-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, N((C<sub>1</sub>-C<sub>6</sub>)-alkyl)<sub>2</sub>, NH(C<sub>1</sub>-C<sub>7</sub>)-acyl, phenyl, or O-(CH<sub>2</sub>)<sub>o</sub>-phenyl,

wherein o is 0-6 and wherein the phenyl ring is optionally substituted one to 3 times, each substituent chosen independently from F, Cl, Br, I, OH, CF<sub>3</sub>, NO<sub>2</sub>, CN, OCF<sub>3</sub>, (C<sub>1</sub>-C<sub>6</sub>)-alkoxy, (C<sub>1</sub>-C<sub>6</sub>)-alkyl, NH<sub>2</sub>, NH(C<sub>1</sub>-C<sub>6</sub>)-alkyl, N((C<sub>1</sub>-C<sub>6</sub>)-alkyl)<sub>2</sub>, SO<sub>2</sub>-CH<sub>3</sub>, COOH, COO-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, and CONH<sub>2</sub>;

A is (C<sub>0</sub>-C<sub>15</sub>)-alkanediyl, wherein one or more carbon atoms in the alkanediyl radical are optionally replaced, independently of one another, by -O-, -(C=O)-, -CH=CH-, -C≡C-, -S-, -CH(OH)-, -CHF-, -CF<sub>2</sub>-, -(S=O)-, -(SO<sub>2</sub>)-, -N((C<sub>1</sub>-C<sub>6</sub>)-alkyl)-, -N((C<sub>1</sub>-C<sub>6</sub>)-alkylphenyl)- or -NH-;

n is a number from 0 to 4;

Cyc1 is a 3- to 7-membered, saturated, partially saturated or unsaturated ring, wherein 1 carbon atom is optionally replaced by O or S;

R3, R4, R5 are, independently of each other, hydrogen, F, Cl, Br, I, OH, NO<sub>2</sub>, CN, COOH, COO(C<sub>1</sub>-C<sub>6</sub>)-alkyl, CO(C<sub>1</sub>-C<sub>4</sub>)-alkyl, CONH<sub>2</sub>,

CONH(C<sub>1</sub>-C<sub>6</sub>)-alkyl, CON[(C<sub>1</sub>-C<sub>6</sub>)-alkyl]<sub>2</sub>, (C<sub>1</sub>-C<sub>8</sub>)-alkyl, (C<sub>2</sub>-C<sub>6</sub>)-alkenyl, (C<sub>2</sub>-C<sub>6</sub>)-alkynyl, (C<sub>1</sub>-C<sub>12</sub>)-alkoxy, HO-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, or (C<sub>1</sub>-C<sub>6</sub>)-alkoxy-(C<sub>1</sub>-C<sub>6</sub>)-alkyl,

wherein one, more than one or all hydrogens in the alkyl and alkoxy radicals are optionally replaced by fluorine;

SO<sub>2</sub>-NH<sub>2</sub>, SO<sub>2</sub>NH(C<sub>1</sub>-C<sub>6</sub>)-alkyl, SO<sub>2</sub>N[(C<sub>1</sub>-C<sub>6</sub>)-alkyl]<sub>2</sub>, S-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, S-(CH<sub>2</sub>)<sub>o</sub>-phenyl, SO-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, SO-(CH<sub>2</sub>)<sub>o</sub>-phenyl, SO<sub>2</sub>-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, or SO<sub>2</sub>-(CH<sub>2</sub>)<sub>o</sub>-phenyl,

wherein o is 0-6 and wherein the phenyl radical is optionally substituted up to twice, each substituent chosen

independently from F, Cl, Br, OH, CF<sub>3</sub>, NO<sub>2</sub>, CN, OCF<sub>3</sub>, (C<sub>1</sub>-C<sub>6</sub>)-alkoxy, (C<sub>1</sub>-C<sub>6</sub>)-alkyl, and NH<sub>2</sub>;

NH<sub>2</sub>, NH-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, N((C<sub>1</sub>-C<sub>6</sub>)-alkyl)<sub>2</sub>, NH(C<sub>1</sub>-C<sub>7</sub>)-acyl, phenyl, (CH<sub>2</sub>)<sub>o</sub>-phenyl, O-(CH<sub>2</sub>)<sub>o</sub>-phenyl,

wherein o is 0-6 and wherein the phenyl ring is optionally substituted one to 3 times, each substituent chosen

independently from F, Cl, Br, I, OH, CF<sub>3</sub>, NO<sub>2</sub>, CN, OCF<sub>3</sub>,

(C<sub>1</sub>-C<sub>8</sub>)-alkoxy, (C<sub>1</sub>-C<sub>6</sub>)-alkyl, NH<sub>2</sub>, NH(C<sub>1</sub>-C<sub>6</sub>)-alkyl,

N((C<sub>1</sub>-C<sub>6</sub>)-alkyl)<sub>2</sub>, SO<sub>2</sub>-CH<sub>3</sub>, COOH, COO-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, and CONH<sub>2</sub>;

or

R3 and R4 together with the carbon atoms carrying them are a 5- to 7-membered, saturated, partially or completely unsaturated ring Cyc2,

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wherein 1 or 2 carbon atoms in the ring are optionally replaced by N, O or S, and

wherein Cyc2 is optionally substituted by (C<sub>1</sub>-C<sub>6</sub>)-alkyl, (C<sub>2</sub>-C<sub>5</sub>)-alkenyl, (C<sub>2</sub>-C<sub>5</sub>)-alkynyl,

wherein, in each substituent of Cyc2, one CH<sub>2</sub> group is optionally replaced by O, or substituted by H, F, Cl, OH, CF<sub>3</sub>, NO<sub>2</sub>, CN, COO(C<sub>1</sub>-C<sub>4</sub>)-alkyl, CONH<sub>2</sub>, CONH(C<sub>1</sub>-C<sub>4</sub>)-alkyl, or OCF<sub>3</sub>, and

R5 is hydrogen;

or a pharmaceutically acceptable salt thereof.

2. A compound as claimed in claim 1, wherein A is linked to the thienyl ring in position 2.
3. A compound as claimed in claim 1, wherein

R1, R2 are, independently of each other, hydrogen, F, Cl, Br, I, OH, NO<sub>2</sub>, CN, COOH, CO(C<sub>1</sub>-C<sub>6</sub>)-alkyl, COO(C<sub>1</sub>-C<sub>6</sub>)-alkyl, CONH<sub>2</sub>, CONH(C<sub>1</sub>-C<sub>6</sub>)-alkyl, CON[(C<sub>1</sub>-C<sub>6</sub>)-alkyl]<sub>2</sub>, (C<sub>1</sub>-C<sub>8</sub>)-alkyl, (C<sub>2</sub>-C<sub>6</sub>)-alkenyl, (C<sub>2</sub>-C<sub>6</sub>)-alkynyl, (C<sub>1</sub>-C<sub>6</sub>)-alkoxy, HO-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, (C<sub>1</sub>-C<sub>6</sub>)-alkoxy-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, phenyl, benzyl, (C<sub>1</sub>-C<sub>4</sub>)-alkylcarbonyl, or SO-(C<sub>1</sub>-C<sub>6</sub>)-alkyl,

wherein one, more than one or all hydrogens in the alkyl or alkoxy radicals are optionally replaced by fluorine;

A is (C<sub>0</sub>-C<sub>15</sub>)-alkanediyl, wherein one or more carbon atoms in the alkanediyl radical are optionally replaced, independently of one another, by -O-, -(C=O)-, -CH=CH-, -C≡C-, -S-, -CH(OH)-, -CHF-, -CF<sub>2</sub>-, -(S=O)-, -(SO<sub>2</sub>)-, -N((C<sub>1</sub>-C<sub>6</sub>)-alkyl)-, -N((C<sub>1</sub>-C<sub>6</sub>)-alkylphenyl)- or -NH-;

n is a number 2 or 3;

Cyc1 is a 5- to 6-membered, saturated, partially saturated or unsaturated ring, wherein 1 carbon atom is optionally replaced by O or S;

R3, R4, R5 are, independently of each other, hydrogen, F, Cl, Br, I, OH, NO<sub>2</sub>, CN, COOH, COO(C<sub>1</sub>-C<sub>6</sub>)-alkyl, CO(C<sub>1</sub>-C<sub>4</sub>)-alkyl, CONH<sub>2</sub>, CONH(C<sub>1</sub>-C<sub>6</sub>)-alkyl, CON[(C<sub>1</sub>-C<sub>6</sub>)-alkyl]<sub>2</sub>, (C<sub>1</sub>-C<sub>8</sub>)-alkyl, (C<sub>2</sub>-C<sub>6</sub>)-alkenyl, (C<sub>2</sub>-C<sub>6</sub>)-alkynyl, (C<sub>1</sub>-C<sub>12</sub>)-alkoxy, HO-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, (C<sub>1</sub>-C<sub>6</sub>)-alkoxy-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, (C<sub>1</sub>-C<sub>4</sub>)-alkylphenyl, (C<sub>1</sub>-C<sub>4</sub>)-alkoxyphenyl, S-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, or SO-(C<sub>1</sub>-C<sub>6</sub>)-alkyl,

wherein one, more than one or all hydrogens in the alkyl or alkoxy radicals are optionally replaced by fluorine;

or

R3 and R4 together with the carbon atoms carrying them are a 5- to 7-membered, saturated, partially or completely unsaturated ring Cyc2,

wherein 1 or 2 carbon atoms in the ring are optionally replaced by N, O or S, and

wherein Cyc2 is optionally substituted by (C<sub>1</sub>-C<sub>6</sub>)-alkyl, (C<sub>2</sub>-C<sub>5</sub>)-alkenyl, or (C<sub>2</sub>-C<sub>5</sub>)-alkynyl,

wherein in each substituent of Cyc2, one CH<sub>2</sub> group is optionally replaced by O, or substituted by H, F, Cl, OH, CF<sub>3</sub>, NO<sub>2</sub>, CN, COO(C<sub>1</sub>-C<sub>4</sub>)-alkyl, CONH<sub>2</sub>, CONH(C<sub>1</sub>-C<sub>4</sub>)-alkyl, or OCF<sub>3</sub>, and

R5 is hydrogen.

4. A compound as claimed in claim 1, wherein

R1, R2 are, independently of each other, hydrogen, (C<sub>1</sub>-C<sub>6</sub>)-alkyl, (C<sub>1</sub>-C<sub>4</sub>)-alkoxy, HO-(C<sub>1</sub>-C<sub>4</sub>)-alkyl, (C<sub>1</sub>-C<sub>4</sub>)-alkoxy-(C<sub>1</sub>-C<sub>4</sub>)-alkyl, F, Cl, CF<sub>3</sub>, OCF<sub>3</sub>, OCH<sub>2</sub>CF<sub>3</sub> (C<sub>1</sub>-C<sub>4</sub>)-alkyl-CF<sub>2</sub>-, phenyl, benzyl, (C<sub>1</sub>-C<sub>4</sub>)-alkylcarbonyl, (C<sub>2</sub>-C<sub>4</sub>)-alkenyl, (C<sub>2</sub>-C<sub>4</sub>)-alkynyl, or COO(C<sub>1</sub>-C<sub>4</sub>)-alkyl;

A is -CH=CH-CH<sub>2</sub>- or (C<sub>1</sub>-C<sub>4</sub>)-alkanediyl, wherein one or two CH<sub>2</sub> groups are optionally replaced by - (C=O)-, -CH=CH-, -CH(OH)-, -NH-, -CHF-, -CF<sub>2</sub>-, or -O-;

n is a number 2 or 3;

Cyc1 is unsaturated ring, wherein 1 carbon atom is optionally replaced by O or S;

R3, R4, R5 are, independently of each other, hydrogen, F, Cl, Br, I, NO<sub>2</sub>, OH, CN, (C<sub>1</sub>-C<sub>6</sub>)-alkyl, (C<sub>1</sub>-C<sub>8</sub>)-alkoxy, OCF<sub>3</sub>, OCH<sub>2</sub>CF<sub>3</sub>, S-(C<sub>1</sub>-C<sub>4</sub>)-

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alkyl, COOH, HO-(C<sub>1</sub>-C<sub>4</sub>)-alkyl, (C<sub>1</sub>-C<sub>4</sub>)-alkoxy-(C<sub>1</sub>-C<sub>4</sub>)-alkyl, (C<sub>1</sub>-C<sub>2</sub>)-alkylphenyl, or (C<sub>1</sub>-C<sub>2</sub>)-alkoxyphenyl, or

R3 and R4 together are -CH=CH-O-, -CH=CH-S-, -O-(CH<sub>2</sub>)<sub>p</sub>-O-, -O-CF<sub>2</sub>-O-, or -CH=CH-CH=CH-, wherein p = 1 or 2, and

R5 is hydrogen.

5. A compound as claimed in claim 1, wherein R2 is hydrogen.

6. A compound as claimed in claim 1, wherein

R1 is hydrogen, CF<sub>3</sub>, (C<sub>1</sub>-C<sub>4</sub>)-alkyl, or phenyl,

R2 is hydrogen,

A is -CH<sub>2</sub>-, -C<sub>2</sub>H<sub>4</sub>-, -C<sub>3</sub>H<sub>6</sub>-, -CH(OH)-, -(C=O)-, -CH=CH-, -CH=CH-CH<sub>2</sub>-, -CO-CH<sub>2</sub>-CH<sub>2</sub>- or -CO-NH-CH<sub>2</sub>-;

n is a number 2 or 3;

Cyc1 is unsaturated ring, wherein 1 carbon atom is optionally replaced by S;

R3,R4,R5 are, independently of each other, hydrogen, F, Cl, I, NO<sub>2</sub>, OH, CN, (C<sub>1</sub>-C<sub>6</sub>)-alkyl, (C<sub>1</sub>-C<sub>8</sub>)-alkoxy, O-CH<sub>2</sub>-phenyl, OCF<sub>3</sub>, S-CH<sub>3</sub>, or COOH or

R3 and R4 together are -CH=CH-O-, -O-(CH<sub>2</sub>)<sub>p</sub>-O-, -O-CF<sub>2</sub>-O-, -CH=CH-CH=CH-, wherein p = 1 or 2, and

R5 is hydrogen.

7. A compound as claimed in claim 1, wherein  
A is -CH<sub>2</sub>- or -CH<sub>2</sub>-CH<sub>2</sub>-.
8. A compound as claimed in claim 1, wherein  
Cyc1 is phenyl.
9. A compound as claimed in claim 1, wherein  
Cyc1 is thienyl.
10. A compound as claimed in claim 1, wherein  
Cyc1 is monosubstituted.
11. A medicament comprising at least one compound as claimed in claim 1.
12. A medicament comprising at least one compound as claimed in claim 1 and at least one more blood glucose-lowering active ingredient.
13. A method for treating type 1 or type 2 diabetes, comprising administering to a patient in need thereof an effective amount of at least one compound as claimed in claim 1.
14. A method for lowering blood glucose, comprising administering to a patient in need thereof an effective amount of at least one compound as claimed in claim 1.

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15. A method for treating type 1 or type 2 diabetes, comprising administering to a patient in need thereof an effective amount of at least one compound as claimed in claim 1 and at least one other active ingredient, wherein the at least one other active ingredient is effective for lowering blood glucose.
16. A method for lowering blood glucose, comprising administering to a patient in need thereof an effective amount of at least one compound as claimed in claim 1 and at least one other active ingredient, wherein the at least one other active ingredient is effective for lowering blood glucose.
17. A process for producing a medicament comprising at least one compound as claimed in claim 1, comprising:  
mixing the at least one compound as claimed in claim 1 with a pharmaceutically suitable carrier, and  
converting this mixture into a form suitable for administration.